## **PCT**

#### WORLD INTELLECTUAL PROPERTY ORGANIZATION International Bureau



#### INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 6:

(11) International Publication Number:

WO 95/16224

G03C 5/08, B41M 3/14, B42D 15/10, G03C 8/28, 8/52

A1

(43) International Publication Date:

15 June 1995 (15.06.95)

(21) International Application Number:

PCT/EP94/04095

(22) International Filing Date:

9 December 1994 (09.12.94)

(81) Designated States: AU, BR, CA, CN, JP, US, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC,

NL, PT, SE).

(30) Priority Data:

93203472.1

10 December 1993 (10.12.93) EP

(34) Countries for which the regional or international application was filed:

DE et al.

(71) Applicants (for all designated States except US): AGFA-GEVAERT NAAMLOZE VENNOOTSCHAP [BE/BE]; Septestraat 27, B-2640 Mortsel (BE). NATIONALE BANK VAN BELGIË N.V. [BE/BE]; De Berlaimontlaan 14, B-1000 Brussel (BE).

(72) Inventors; and

(75) Inventors/Applicants (for US only): VERMEULEN, Leo [BE/BE]; Agfa-Gevaert N.V., DIE 3800, Septestraat 27, B-2640 Mortsel (BE). DE BAETS, Daniel [BE/BE]; Nationale Bank van België N.V., De Berlaimontlaan 14, B-1000 Brussel (BE).

**Published** 

With international search report.

Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.

(54) Title: OPAQUE DOCUMENT CONTAINING INTERFERENCE PIGMENTS PROVIDING EASY VERIFICATION AND PRO-TECTION AGAINST PHOTO-COPYING

#### (57) Abstract

A security document containing a photograph obtained by the silver complex or dye diffusion transfer process in an image receiving layer containing light interference pigments uniformly distributed in said image receiving layer, and/or in a layer overlaying or underlying said image receiving layer, said layer(s) being present on an opaque support, characterised in that said interference pigments are composed of silicate platelets coated with either. A) a first layer of highly refractive metal oxide, and B) a second black layer essentially consisting of metallic iron, molybdenum and/or wolfram, or coated with: A) a first layer of highly refractive metal oxide, and B) a second black layer essentially consisting of carbon or metal, and C) a third layer of highly refractive metal oxide.

## FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

ΑT	Austria	GB	United Kingdom	MR	Mauritania
AU	Australia	GE	Georgia	MW	Malawi
BB	Barbados	GN	Guinea	NE	Niger
BE	Belgium	GR	Greece	NL	Netherlands
BF	Burkina Faso	HU	Hungary	NO	Norway
BG	Bulgaria	<b>IE</b>	Ireland	NZ	New Zealand
BJ	Benin	<b>FT</b>	Italy	PL	Poland
BR	Brazil	JP	Japan	PT	Portugal
BY	Belarus	KE	Kenya	RO	Romania
CA	Canada	KG	Kyrgystan	RU	Russian Federation
CF	Central African Republic	KР	Democratic People's Republic	SD	Sudan
CG	Congo		of Korea	SE	Sweden
CH	Switzerland	KR	Republic of Korea	SI	Slovenia
CI	Côte d'Ivoire	KZ	Kazakhstan	SK	Slovakia
CM	Cameroon	LI	Liechtenstein	SN	Senegal
CN	China	LK	Sri Lanka	TD	Chad
CS	Czechoslovakia	LU	Luxembourg	TG	Togo
CZ	Czech Republic	LV	Latvia	TJ	Tajikistan
DE	Germany	MC	Monaco	TT	Trinidad and Tobago
DK	Denmark	MD	Republic of Moldova	UA	Ukraine
ES	Spain	MG	Madagascar	US	United States of America
FI	Finland	ML	Mali	UZ	Uzbekistan
FR	Prance	MN	Mongolia	VN	Viet Nam
GA	Gabon				

WO 95/16224

. 1

PCT/EP94/04095

OPAQUE DOCUMENT CONTAINING INTERFERENCE PIGMENTS PROVIDING EASY VERIFICATION AND PROTECTION AGAINST PHOTO-COPYING

#### 1. Field of the Invention

The present invention relates to opaque security documents that can be verified on their authenticity and are protected against counterfeiting by photo-copying.

#### 2. Background of the Invention

Security documents that must be verifiable on their authenticity are e.g. all kinds of identification documents such as passports, visas, identity cards, driver licenses, bank cards, credit cards, security entrance cards, and further value-documents such as banknotes, shares, bonds, certificates, cheques, lottery tickets and all kinds of entrance tickets such as airplane tickets and railroad season-tickets.

Nowadays, by the availability of markedly improved black-andwhite and color copiers it becomes more and more easy to copy documents at high quality hardly to distinguish from the originals.

To protect the above mentioned documents against fraudulent alterations and reproduction by photo-copying different techniques are used such as the melt-laminating or glueing thereto of preprinted plastic overlayers; the printing with special inks; the application of coatings or layers for example loaded with magnetic or fluorescent pigments; coloring or metallizing the substrate of the documents; incorporating holograms; applying fine line printing, watermarks, fibers, security threads, light diffraction marks, liquid crystal marks and/or substances called nacreous, iridiscent or interference pigments.

In a particular case disclosed in US-P 4.151,666 light-transmissive pigments serving as diffuse reflectors are applied by printing to form a verification pattern in a laminated identification card (I.D. card). In the specification of the same US-P the use of nacreous pigments in verification patterns has been described. Nacreous pigments, also called pearlescent pigments have light-reflection characteristics that change as a function of the viewing or copying angle. The effect of changing color with viewing angle makes that nacreous pigments represent a simple and convenient matter to built in a verification feature associated with a non-

copyable optical property.

## 3. Objects and Summary of the Invention

It is an object of the present invention to provide opaque security documents incorporating a photograph and containing a verification feature that can not be copied by photographic techniques and wherein there is no need for specialized devices or conditions to verify the document on its authenticity.

It is more particularly an object of the present invention to provide opaque security documents containing at least one photograph obtained by means of the silver complex or dye diffusion transfer process in combination with interference pigments applied by coating before the photograph is formed.

It is a special object of the present invention to provide an opaque security document containing a portrait in combination with a plurality of interference pigments of different color for easy verfication by the naked eye of the security document involved.

Other objects and advantages of the present invention will become clear from the further description and examples.

In accordance with the present invention there is provided a document containing a photograph obtained by the silver complex or dye diffusion transfer process in an image receiving layer. characterized in that said document contains light interference pigments uniformly distributed in said image receiving layer, and/or in a layer overlaying or underlying said image receiving layer, said layer(s) being present on an opaque support.

By "opaque support" in the document according to the present invention has to be understood a support having a visible light-blocking capacity of at least 70 %, not excluding supports that have inherently a color or contain colorants, e.g. dyes or pigments.

Said layers containing said uniformly distributed interference pigments may be present at both sides of said opaque support.

## 4. Detailed Description of the Invention

Nacreous or pearlescent pigments being interference pigments were originally named for their simulation of pearly luster.

Interference pigments are in the form of light-reflecting crystal platelets of appropriate thickness to produce color by

WO 95/16224

3

PCT/EP94/04095

interference. These pigments exhibit a color play that verges on iridiscence and under a given angle of reflection will allow only the copying of a single color, whereas other colors appear under different angles of reflection, in other words these pigments show another color to the human eye depending on the observation angle.

Very good anticopying results are obtained with synthetic inorganic interference pigments chief among which are titanium dioxide-coated mica or other metal-oxide coated pigments in which the metal oxide has preferably a refractive index comparable with the refractive index of TiO<sub>2</sub>, e.g. ZrO<sub>2</sub>, Fe<sub>2</sub>O<sub>3</sub> or Cr<sub>2</sub>O<sub>3</sub>.

The platelets of metal oxide coated mica pigments have three layers in such a way that on each of the broad faces of the mica platelets a very thin coating of metal oxide is present.

The more brilliant interference pigments approach metallic luster, an effect that is enhanced by the presence of absorption colorants.  ${\rm TiO_2}$ -mica readily lends itself to incorporation of absorption colorants. Ferric oxide (Fe<sub>2</sub>O<sub>3</sub>) added to the  ${\rm TiO_2}$  layer, for example imparts a yellow color which in conjunction with a yellow interference color creates gold. When  ${\rm Fe_2O_3}$  is used in place of  ${\rm TiO_2}$  as the oxide coating on mica, these pigments have a yellow-red absorption color because of the inherent color of the  ${\rm Fe_2O_3}$ . They range from bronze to deep copper-red and have a metallic luster.

The preparation of such kind of pigments is described in "Nacreous (Pearlescent) Pigments and Interference Pigments by L.M. Greenstein Henry L. Mattin Laboratories Reprinted from Pigment Handbook, Vol. I. Properties and Economics, Second Edition, Edited by Peter A. Lewis, (1988) by permission of John Wiley & Sons, Inc. The Mearl Corporation, 41 East 42nd Street, New York, N.Y. 100017.

Mica pigments serving as a substrate of the interference coatings are a group of hydrous aluminum silicate minerals with platy morphology and perfect basal (micaceous) cleavage. Examples of suitable micas are e.g. muscovite  $KAl_2(AlSi_3O_{10})(OH)_2$ , paragonite  $NaAl_2(AlSi_3O_{10})(OH)_2$ , phlogopite  $K(Mg,Fe)(AlSi_3O_{10})(OH)_2$ , biotite  $K(Fe,Mg)(AlSi_3O_{10})(OH)_2$  and lepidolite  $K(Li,Al)_{2.5-3.0}(Al_{1.0}-0.5Si_3.0-3.5O_{10})(OH)_2$  etc..

Details about the application of metal and/or metal oxide coatings on the mica platelets are further given in US-P 3,087,827, 4,434,010 and 5,059,245 and in published EP-A-45 851, EP-A-313 280, DE-A-11 65 182, DE-A-32 37 264, DE-A-38 25 702 and DE-A-36 17 430

PCT/EP94/04095 WO 95/16224

4

and DE-OS 41 41 069.

In a preferred embodiment according to the present invention interference pigments having a composition as described in published German patent application DE-OS 41 41 069 are used. Said interference pigments, called "Glanzpigmente" according to said DE-OS are composed of silicate platelets coated with either:

- A) a first layer of highly refractive metal oxide, and
- B) a second black layer essentially consisting of metallic iron, molybdenum and/or wolfram, or coated with :
- A) a first layer of highly refractive metal oxide, and
- B) a second black layer essentially consisting of carbon or metal, and
- C) a third layer of highly refractive metal oxide. By the presence of a semi-transparent carbon layer said platelets have a high chemical resistance. The carbon layer is obtained by thermal decomposition of oxygen-containing hydrocarbon compounds which for each two carbon atoms contain at least one oxygen atom e.g. as in carbohydrates such as sorbitol.

The metal layer B) can be formed in a medium of inert gas from ingas-phase-decomposable metal compounds, e.g. metal carbonyl compounds, and the metal oxide layers A) and C) are formed by decomposition in gas phase of volatile metal compounds in the presence of oxygen water vapour or mixtures thereof.

The first layer A) consists e.g. of the oxides of titanium, zirconium, tin and/or iron.

Mica platelets double-side coated with one or more metal oxide layers for use as interference pigments are commercially available e.g. under the tradenames IRIODINE (E. Merck, Darmstadt), FLONAC (Kemira Oy, Pori, Finland), MEARLIN (The Mearl Corporation, New York, U.S.A.) and PALIOSECURE (BASF, Germany). Under these tradenames interference pigments showing violet, red, green and blue colors are available on the market.

Preferred interference platelet-type pigments for use according to the present invention have a largest surface diameter preferably between 5 and 200  $\mu m$  and more preferably of 25  $\mu m$  to 30  $\mu m$ . The thickness of the platelet-type interference pigments is preferably between 0.1  $\mu$ m and 0.6  $\mu$ m and more preferably between 0.2  $\mu$ m and 0.4 μm.

In accordance with a preferred embodiment particularly good anti-copying security is obtained by means of an opaque document

5

wherein said uniformly distributed interference pigments of a particular color are present in combination with a printed pattern containing interference pigments of a color different from the color of the uniformly distributed interference pigments when viewed under the angle of reflection under which photocopying could take place.

The "copying angle" is the angle, usually 90°, formed by the information-containing light rays and the copying material.

In accordance with another embodiment there is provided an opaque document wherein said uniformly distributed interference pigments are present in conjunction with different and pattern-wise printed interference pigments on a background having a color substantially the same as the color, e.g. yellow, reflected by said pattern-wise printed interference pigments under the angle of reflection under which photocopying could take place. Hereby the pattern of the pattern-wise printed interference pigments becomes undetectable against the image background for the copying machine, but is still visible in the document under a viewing angle different from the copying angle.

In accordance with a further embodiment there is provided an opaque document wherein said uniformly distributed interference pigments are present in combination with pattern-wise printed colored light absorbing substances (colored pigments or dyes), and/or white light-diffusing pigments.

The layer containing the uniformly distributed interference pigments may be applied by coating techniques operating with a coating liquid containing said pigments in dispersed form and a dissolved binding agent or containing said pigments dispersed together with a binding agent in the form of a latex. After coating the solvent or dispersing liquid, e.g. water, is removed by evaporation. Any coating technique for the application of thin liquid layers may be used as is known e.g. from the field of the manufacture of photographic silver halide emulsion layer materials, e.g. doctor blade coating, gravure roller coating, meniscus coating, air knife coating, slide hopper coating and spraying.

According to a special coating technique the interference pigments are applied in a radiation-curable binder or binder system wherein e.g. monomers act as solvent for polymers or prepolymers as described e.g. in published EP-A 0 522 609, so that after coating of the liquid coating composition no solvent has to be evaporated.

In accordance with a particular embodiment the uniformly

6

distributed interference pigments are applied in a layer that is transferred by a stripping-off procedure to built the document of the present invention. Such procedure is described in published EP-A 0 478 790 but is applied therein for controlling the whiteness of an image present on a permanent support using for the stripping-off and transfer procedure a temporary support coated with a wetstrippable non-photosensitive layer containing fluorescent whitening agent(s) in a hydrophilic colloid binder.

According to a special embodiment the interference pigments applied in the form of a pigment-transfer-foil wherefrom by hot transfer the pigments are transferred uniformly onto the substrate of the security document.

Still another coating technique suited for uniformly applying said pigments is by dry powder-spraying optionally on a hot-melt resin layer wherein they are impregnated by pressure and heat. On top of the pigments an adhesive, e.g. wax may be applied to improve adherence to the selected substrate. That substrate may have hydrophilic or hydrophobic surface properties.

Spray-coating may be applied for covering the whole surface of the substrate or only a part thereof producing "light interference rainbow-effects". By using varying mixtures of different interference pigments the intensity of one color can be made to decrease gradually while an increasing color intensity of another pigment comes up. The human eye will see the rainbow effect varying according to the perception angle and will recognize the basic color of each of the sprayed pigments, but a photocopier operating with a fixed copying angle will only reproduce, say a single yellowish-brown color and not the colors of the interference pigments that can be seen under different inspection angles.

The interference pigments can be used for pigmenting a commercial coating varnish which may then be used for pre-coating a security document substrate, e.g. opaque resin film or paper. The coating may proceed with common varnishing or impregnation machinery instead of using printing presses.

As already mentioned herein the uniformly applied interference pigments are advantageously combined with image-wise or pattern-wise interference pigments of another color.

The image-wise or pattern-wise application of interference pigments proceeds by printing with an ink containing said pigments. Suited printing processes are e.g. planographic offset printing,

7

gravure printing, intaglio printing, screen printing, flexographic printing, relief printing, tampon printing, ink jet printing and toner-transfer printing from electro(photo)graphic recording materials.

For use in printing on hydrophilic layers or substrates the ink contains for example a 15 to 20 % by weight mixture of the interference pigments in a solution of cellulose nitrate in a polyethylene ether. Such ink has a good adherence on hydrophilic colloid layers such as gelatin-containing layers used in DTR-recording materials. Said ink is advantageously applied with a commercial screen press using a polyester screen with a 77 and 55 mesh. The interference colors gradually appear on drying the ink.

Thus applied ink patterns on a hydrophilic image-receiving layer for DTR-image production remain unchanged during DTR-processing.

The presence of the interference pigments in one of the layers of the opaque security document does not affect the possibility to print thereon further graphic or alpha-numerical information by any known printing technique.

For easy visual verification the interference pigments are present preferably in a coverage of  $0.3~\rm g/m^2$  to  $10~\rm g/m^2$  and more preferably in a coverage between  $0.7~\rm g/m^2$  and  $3~\rm g/m^2$  in the above mentioned silver-type DTR and dye diffusion transfer materials, either in the image-receiving layer itself and/or in a waterpermeable top layer and/or in a subbing layer covering the support.

The presence of a dried water-impermeable ink pattern on the image-receiving layer blocks DTR-image formation. Thereby it is possible to arrange e.g. fine line patterns such as guilloches in the photograph creating that way an additional security feature.

The printing of an interference pigment-containing pattern may proceed on a substrate already covered e.g. by a hologram, light-diffraction pattern, metallic pattern that can be viewed throught the printed pattern so that the properties of the interference pigments are added thereto.

The printed pattern containing interference pigments forms no obstacle for a good adherence with laminated plastic resinous covering material. By proper selection of the binder of the ink it can be co-melted with the resin material laminated thereto.

According to a particular embodiment the interference pigmentcontaining ink is applied on a temporary support, e.g. polystyrene

8

support, wherefrom the ink layer can be stripped off and transferred to a permanent support, e.g. a glued and preprinted substrate of a security document. The ink layer, applied overall or pattern-wise, after leaving the temporary support covers underlying pre-printed data on the permanent support. For preventing fraudulent copying these data have the same color as the interference pigment layer when seen under the copying angle. Insufficient image contrast is available so that copying of the pre-printed data is no longer possible. By applying a dried interference pigment-containing ink layer that is translucent the underlying data can be visually inspected therethrough by altering the perception angle.

In accordance with the preceding embodiment a security document according to the present invention, e.g. serving as I.D. card, is preferably in the form of a laminate in which the information-containing layer(s) are sealed between protective resinous sheets. I.D. card laminates may be built up as described e.g. in US-P 4.101.701, US-P 4.762,759, US-P 4.902.593, published EP-A 0 348 310 and published EP-A 0 462 330. By lamination tamper-proof documents are produced which do not allow the opening of the laminate without damaging the image contained therein. The destruction of the seal will leave visual fraude traces on the security document.

In accordance with a first mode in the opaque security document according to the present invention a black-and-white photograph in the form of a silver image is formed by the silver salt diffusion transfer process, called herein DTR-process. According to said process dissolved silver halide salt is transferred imagewise in a special image receiving layer, called development nuclei containing layer, for reducing therein transferred silver salt, said development nuclei containing layer contains itself and/or in an overlaying and/or an underlaying layer uniformly distributed therein said interference pigments.

The principles of the DTR-process are described in U.S. patent No. 2,352,014 of André Rott, issued June 20, 1944. According to said process silver complexes are image-wise transferred by diffusion from a silver halide emulsion layer to an image-receiving layer, where they are converted, in the presence of development nuclei, into a silver image. For this purpose, an image-wise exposed silver halide emulsion layer is developed by means of a developing substance in the presence of a so-called silver halide solvent. In

WO 95/16224

the exposed parts of the silver halide emulsion layer the silver halide is developed to metallic silver so that it cannot dissolve anymore and consequently cannot diffuse. In the non-exposed parts of the silver halide emulsion layer the silver halide is converted into soluble silver complexes by means of a silver halide complexing agent, acting as silver halide solvent, and said complexes are transferred by diffusion into an image-receiving layer being in waterpermeable contact with said emulsion layer to form by the catalytic action of said development nuclei, in so-called physical development, a silver-containing image in the image-receiving layer.

9

PCT/EP94/04095

More details on the DTR-process can be found in "Photographic Silver Halide Diffusion Processes" by A. Rott and E. Weyde, Focal Press, London, New York (1972).

In accordance with a second mode in the opaque security document according to the present invention a color photograph in the form of one or more dye images is formed by the dye diffusion transfer process (dye DTR-process) wherein the image-wise transfer of dye(s) is controlled by the development of (a) photo-exposed silver halide emulsion layer(s), and wherein dye(s) is (are) transferred imagewise in a special image receiving layer, called mordant layer, for fixing the dyes, said mordant layer and/or an overlaying and/or an underlaying layer containing uniformly distributed therethrough said interference pigments.

Dye diffusion transfer reversal processes are based on the image-wise transfer of diffusible dye molecules from an image-wise exposed silver halide emulsion material into a waterpermeable image-receiving layer containing a mordant for the dye(s). The image-wise diffusion of the dye(s) is controlled by the development of one or more image-wise exposed silver halide emulsion layers, that for the production of a multicolor image are differently spectrally sensitized and contain respectively a yellow, magenta and cyan dye molecules. A survey of dye diffusion transfer imaging processes has been given by Christian C. Van de Sande in Angew. Chem. - Ed. Engl. 22 (1983) n° 3, 191-209 and a particularly useful process is described in US-P 4.496,645.

For use in dye diffusion transfer photography the type of mordant chosen will depend upon the dye to be mordanted. If acid dyes are to be mordanted, the image-receiving layer being a dyemordanting layer contains basic polymeric mordants such as polymers of amino-guanidine derivatives of vinyl methyl ketone such as

10

described in US-P 2.882,156, and basic polymeric mordants and derivatives, e.g. poly-4-vinylpyridine, the metho-p-toluene sulphonate of poly-2-vinylpyridine and similar compounds described in US-P 2,484,430, and the compounds described in the published DE-A 2,009,498 and 2,200,063. Other mordants are long-chain quaternary ammonium or phosphonium compounds or ternary sulphonium compounds, e.g. those described in US-P 3,271,147 and 3,271,148,, and cetyltrimethyl-ammonium bromide. Certain metal salts and their hydroxides that form sparingly soluble compounds with the acid dyes may be used too. The dye mordants are dispersed or molecularly divided in one of the usual hydrophilic binders in the image-receiving layer, e.g. in gelatin, polyvinylpyrrolidone or partly or completely hydrolysed cellulose esters.

In US-P 4,186,014 cationic polymeric mordants are described that are particularly suited for fixing anionic dyes, e.g. sulphinic acid salt dyes that are image-wise released by a redox-reaction described e.g. in in published EP-A 0,004,399 and US-P 4,232,107.

The DTR process can be utilized for reproducing line originals e.g. printed documents, as well as for reproducing continuous tone originals, e.g. portraits.

By the fact that the DTR-image is based on diffusion transfer of imaging ingredients the image-receiving layer and optionally present covering layer(s) have to be waterpermeable.

The reproduction of black-and-white continuous tone images by the DTR-process requires the use of a recording material capable of yielding images with considerable lower gradation than is normally applied in document reproduction to ensure the correct tone rendering of continuous tones of the original. In document reproduction silver halide emulsion materials are used which normally mainly contain silver chloride. Silver chloride not only leads to a more rapid development but also to high contrast.

In U.S. patent. No. 3,985,561, to be read in conjunction herewith, a light-sensitive silver halide material is described wherein the silver halide is predominantly chloride and this material is capable of forming a continuous tone image on or in an image-receiving material by the diffusion transfer process.

According to said U.S. patent a continuous tone image is produced by the diffusion transfer process in or on an image-receiving layer through the use of a light-sensitive layer which

WO 95/16224

contains a mixture of silver chloride and silver iodide and/or silver bromide dispersed in a hydrophilic colloid binder e.g. gelatin, wherein the silver chloride is present in an amount of at least 90 mole % based on the total mole of silver halide and wherein the weight ratio of hydrophilic colloid to silver halide, expressed as silver nitrate, is between 3:1 and about 10:1 by weight.

11

PCT/EP94/04095

With these light-sensitive materials successful reproduction of continuous tone images can be obtained probably as a result of the presence of the indicated amounts of silver iodide and/or silver bromide and of the defined high ratio of hydrophillic colloid to silver halide.

According to U.S. patent No. 4,242,436 likewise to be read in conjunction herewith, the reproduction of continuous tone images can be improved by developing the photographic material with a mixture of developing agents comprising an o-dihydroxybenzene, e.g. catechol, a 3-pyrazolidinone e.g. a 1-aryl-3-pyrazolidinone and optionally a p-dihydroxybenzene, e.g. hydroquinone, the molar amount of the o-dihydroxybenzene in said mixture being larger than the molar amount of the 3-pyrazolidinone, and the p-dihydroxybenzene if any being present in a molar ratio of at most 5 % with respect to the o-dihydroxybenzene.

Suitable development nuclei for use in the above mentioned physical development in the image receiving layer are e.g. noble metal nuclei e.g. silver, palladium, gold, platinum, sulphides, selenides or tellurides of heavy metals such as Pd, Ag, Ni and Co. Preferably used development nuclei are colloidal PdS, Ag<sub>2</sub>S or mixed silver-nickelsulphide particles. The amount of nuclei used in the image receiving layer is preferably between 0.02 mg/m<sup>2</sup> and 10 mg/m<sup>2</sup>.

The image receiving layer comprises for best imaging results the physical development nuclei in the presence of a protective hydrophilic colloid, e.g. gelatin and/or colloidal silica, polyvinyl alcohol etc..

The transfer behaviour of the complexed silver largely depends on the thickness of the image-receiving layer and the kind of binding agent or mixture of binding agents used in the nuclei containing layer. In order to obtain a sharp image with high spectral density the reduction of the silver salts diffusing into the image receiving layer must take place rapidly before lateral diffusion becomes substantial. An image-receiving material satisfying said purpose is described in US-P 4.859,566.

PCT/EP94/04095

12

An image-receiving material of this type is very suitable for use in connection with the present invention and contains a water-impermeable support coated with (1) an image-receiving layer containing physical development nuclei and interference pigments dispersed in a waterpermeable binder and (2) a waterpermeable top layer free from development nuclei and containing a hydrophilic colloid, in such a way that:

- (i) the total solids coverage of said two layers (1) and (2) is e.g. at most  $2 \text{ g/m}^2$ .
- (ii) in layer (1) the coverage of the nuclei is in the range of 0.1  $mg/m^2$  to 10  $mg/m^2$ , and the coverage of binder is in the range of 0.4 to 1.5  $g/m^2$ , and
- (iii) in said top layer (2) the coverage of hydrophilic colloid is in the range of 0.1 to 0.9  $g/m^2$ .

The coating of said layers proceeds preferably with slide hopper coater or curtain coater known to those skilled in the art.

According to a particular embodiment the nuclei containing layer (1) is present on a nuclei-free underlying hydrophilic colloid undercoat layer or undercoat layer system having a coverage in the range of 0.1 to 1  $g/m^2$  of hydrophilic colloid, the total solids coverage of layers (1) and (2) together with the undercoat being at most 2  $g/m^2$ . In connection with this embodiment the nacreous pigments may be also be included in the undercoat layer or may be included therein instead of being present in the nuclei containing layer.

The undercoat optionally incorporates substances that improve the image quality, e.g. incorporates a substance improving the image-tone or the whiteness of the image background. For example, the undercoat may contain a fluorescent substance, silver complexing agent(s) and/or development inhibitor releasing compounds known for improving image sharpness.

According to a special embodiment the image-receiving layer (1) is applied on an undercoat playing the role of a timing layer in association with an acidic layer serving for the neutralization of alkali of the image-receiving layer. By the timing layer the time before neutralization occurs is established, at least in part, by the time it takes for the alkaline processing composition to penetrate through the timing layer. Materials suitable for neutralizing layers and timing layers are disclosed in Research Disclosure July 1974, item 12331 and July 1975, item 13525.

WO 95/16224

13

PCT/EP94/04095

In the image-receiving layer (1) and/or in said top layer (2) and/or in an alkali-neutralizing undercoat gelatin is used preferably as hydrophilic colloid. In layer (1) gelatin is present preferably for at least 60 % by weight and is optionally used in conjunction with an other hydrophilic colloid, e.g. polyvinyl alcohol, cellulose derivatives, preferably carboxymethyl cellulose, dextran, gallactomannans, alginic acid derivatives, e.g. alginic acid sodium salt and/or watersoluble polyacrylamides. Said other hydrophilic colloid may be used also in the top layer for at most 10 % by weight and in the undercoat in an amount lower than the gelatin content.

The image-receiving layer and/or a hydrophilic colloid layer in water-permeable relationship therewith may comprise a silver halide developing agent and/or silver halide solvent, e.g. sodium thiosulphate in an amount of approximately 0.1 g to approximately 4 g per  $m^2$ .

The image-receiving layer or a hydrophilic colloid layer in water-permeable relationship therewith may comprise colloidal silica.

The image-receiving layer may contain as physical development accelerators, in operative contact with the developing nuclei, thioether compounds such as those described e.g. in DE-A-1,124,354; US-P 4,013,471; US-P 4,072,526 and in EP 26520.

According to a preferred embodiment the processing liquid and/or the DTR image-receiving material contains at least one image toning agent. In said case the image toning agent(s) may gradually transfer by diffusion from said image-receiving material into the processing liquid and keep therein the concentration of said agents almost steady. In practice such can be realized by using the silver image toning agents in a coverage in the range from 1 mg/m $^2$  to 20 mg/m $^2$  in a hydrophilic waterpermeable colloid layer.

A survey of suitable toning agents is given in the above mentioned book of André Rott and Edith Weyde, p. 61-65, preference being given to 1-phenyl-1H-tetrazole-5-thiol, also called 1-phenyl-5-mercapto-tetrazole, tautomeric structures and derivatives thereof such as 1-(2,3-dimethylphenyl)-5-mercapto-tetrazole,

- 1-(3,4-dimethylcyclohexyl)-5-mercapto-tetrazole,
- 1-(4-methylphenyl)-5-mercapto-tetrazole,
- 1-(3-chloro-4-methylphenyl)-5-mercapto-tetrazole,
- 1-(3,4-dichlorophenyl)-5-mercapto-tetrazole. Further particularly

14

useful toning agents are of the class of thiohydantoins and of the class of phenyl substituted mercapto-triazoles. Still further toning agents suitable for use in accordance with the preferred embodiment of the present invention are the toning agents described in published European patent applications 218752, 208346, 218753 and US-P 4,683,189.

In the security documents according to the present invention the opaque support is e.g. an opaque paper support or resin coated paper supports, e.g. polyolefin coated paper and polyethylene coated paper of which the polyethylene layer may contain the interference pigments or other opacity providing pigments such as white  $TiO_2$  particles as described e.g. in published European patent application (EP-A) 0 324 192. The opaque supports reflecting white light may contain optical brightening agents.

Other opaque supports that may be used are resin supports containing in their resin mass dispersed white pigments, e.g.  ${\rm TiO}_2$ , or are such resin supports that contain said pigments dispersed in the resin mass in the presence of light-straying microvoids as described e.g. in EP 349152. Organic resins used for manufacturing said supports, e.g. by extrusion, are polycarbonate, polyester, preferably polyethylene terephthalate, polystyrene and homo- and copolymers of vinyl chloride. Further are mentioned cellulose esters e.g. cellulose triacetate.

The above mentioned DTR image-receiving materials may be used in conjunction with any type of photosensitive material containing a silver halide emulsion layer. For continuous tone reproduction the silver halide comprises preferably a mixture of silver chloride, and silver iodide and/or silver bromide, at least 90 mole % based on the total mole of the silver halide being silver chloride, and the ratio by weight of hydrophillic colloid to silver halide expressed as silver nitrate is between 3:1 and 10:1.

The binder for the silver halide emulsion layer and other optional layers contained on the imaging element is preferably gelatin. But instead of or together with gelatin, use can be made of one or more other natural and/or synthetic hydrophilic colloids, e.g. albumin, casein, zein, polyvinyl alcohol, alginic acids or salts thereof, cellulose derivatives such as carboxymethyl cellulose, modified gelatin, e.g. phthaloyl gelatin etc. The weight ratio in the silver halide emulsion layer of hydrophilic colloid binder to silver halide expressed as equivalent amount of silver

nitrate to binder is e.g. in the range of 1:1 to 10:1, but preferably for continuous tone reproduction is between 3.5:1 and 6.7:1.

The silver halide emulsions may be coarse or fine grain and can be prepared by any of the well known procedures e.g. single jet emulsions, double jet emulsions such as Lippmann emulsions, ammoniacal emulsions, thiocyanate- or thioether-ripened emulsions such as those described in US-P 2,222,264, 3,320,069, and 3,271,157. Surface image emulsions may be used or internal image emulsions may be used such as those described in US-P 2,592,250, 3,206,313, and 3,447,927. If desired, mixtures of surface and internal image emulsions may be used as described in US-P 2,996,382.

The silver halide particles of the photographic emulsions may have a regular crystalline form such as cubic or octahedral form or they may have a transition form. Regular-grain emulsions are described e.g. in J. Photogr. Sci., Vol. 12, No. 5, Sept./Oct. 1964, pp. 242-251. The silver halide grains may also have an almost spherical form or they may have a tabular form (so-called T-grains), or may have composite crystal forms comprising a mixture of regular and irregular crystalline forms. The silver halide grains may have a multilayered structure having a core and shell of different halide composition. Besides having a differently composed core and shell the silver halide grains may comprise also different halide compositions and metal dopants inbetween.

The average size expressed as the average diameter of the silver halide grains may range from 0.2 to 1.2 um, preferably between 0.2 $\mu$ m and 0.8 $\mu$ m, and most preferably between 0.3 $\mu$ m and 0.6 $\mu$ m. The size distribution can be homodisperse or heterodispere. A homodisperse size distribution is obtained when 95 % of the grains have a size that does not deviate more than 30 % from the average grain size.

The emulsions can be chemically sensitized e.g. by adding sulphur-containing compounds during the chemical ripening stage e.g. allyl isothiocyanate, allyl thiourea, and sodium thiosulphate. Also reducing agents e.g. the tin compounds described in BE-A 493,464 and 568,687, and polyamines such as diethylene triamine or derivatives of aminomethane-sulphonic acid can be used as chemical sensitizers. Other suitable chemical sensitizers are noble metals and noble metal compounds such as gold, platinum, palladium, iridium, ruthenium and rhodium. This method of chemical sensitization has been described in the article of R.KOSLOWSKY, Z. Wiss. Photogr. Photophys.

16

PCT/EP94/04095

Photochem. 46, 65-72 (1951).

WO 95/16224

The emulsions can also be sensitized with polyalkylene oxide derivatives, e.g. with polyethylene oxide having a molecular weight of 1000 to 20,000, or with condensation products of alkylene oxides and aliphatic alcohols, glycols, cyclic dehydration products of hexitols, alkyl-substituted phenols, aliphatic carboxylic acids, aliphatic amines, aliphatic diamines and amides. The condensation products have a molecular weight of at least 700, preferably of more than 1000. It is also possible to combine these sensitizers with each other as described in BE-P 537,278 and GB-P 727,982.

The silver halide emulsion may be sensitized panchromatically to ensure reproduction of all colors of the visible part of the spectrum or it may be orthochromatically sensitized.

The spectral photosensitivity of the silver halide can be adjusted by proper spectral sensitization by means of the usual mono- or polymethine dyes such as acidic or basic cyanines, hemicyanines, oxonols, hemioxonols, styryl dyes or others, also trior polynuclear methine dyes e.g. rhodacyanines or neocyanines. Such spectral sensitizers have been described by e.g. F.M. HAMER in "The Cyanine Dyes and Related Compounds" (1964) Interscience Publishers, John Wiley & Sons, New York.

The silver halide emulsions may contain the usual stabilizers e.g. homopolar or salt-like compounds of mercury with aromatic or heterocyclic rings such as mercaptotriazoles, simple mercury salts, sulphonium mercury double salts and other mercury compounds. Other suitable stabilizers are azaindenes, preferably tetra- or penta-azaindenes, especially those substituted with hydroxy or amino groups. Compounds of this kind have been described by BIRR in Z. Wiss. Photogr. Photophys. Photochem. 47, 2-27 (1952). Other suitable stabilizers are i.a. heterocyclic mercapto compounds e.g. phenylmercaptotetrazole, quaternary benzothiazole derivatives, and benzotriazole.

A survey of photographic silver halide emulsions and their preparation is given in Research Disclosure December 1989, item 308119.

Processing of the image-wise exposed photographic silver halide emulsion layer proceeds whilst in contact with an image receiving material according to the invention and is accomplished using an alkaline processing liquid having a pH preferably between 9 and 13. The pH of the alkaline processing liquid may be established using

17

various alkaline substances. Suitable alkaline substances are inorganic alkali e.g. sodium hydroxide, potassium carbonate or alkanolamines or mixtures thereof. Preferably used alkanolamines are tertiary alkanolamines e.g. those described in EP-A-397925, EP-A-397926, EP-A-397927, EP-A-398435 and US-P 4,632,896. A combination of alkanolamines having both a pka above or below 9 or a combination of alkanolamines whereof at least one has a pka above 9 and another having a pka of 9 or less may also be used as disclosed in the Japanese patent applications laid open to the public numbers 73949/61, 73953/61, 169841/61, 212670/60, 73950/61, 73952/61, 102644/61, 226647/63, 229453/63, US-P-4,362,811, US-P 4,568,634 etc.. The concentration of these alkanolamines is preferably from 0.1 mol/1 to 0.9 mol/1.

Suitable developing agents for the exposed silver halide are e.g. hydroquinone-type and 1-pheny1-3-pyrazolidone-type developing agents as well as p-monomethylaminophenol and derivatives thereof. Preferably used is a combination of a hydroquinone-type and 1-pheny1-3-pyrazolidone-type developing agent wherein the latter is preferably incorporated in one of the layers comprised on the support of the photographic material. A preferred class of 1-pheny1-3-pyrazolidone-type developing agents is disclosed in the published EP-A 449340.

According to a preferred embodiment for continuous tone reproduction a mixture of developing agents comprising an odihydroxybenzene, e.g. catechol, a 3-pyrazolidinone e.g. a 1-aryl-3-pyrazolidinone and optionally a p-dihydroxybenzene, e.g. hydroquinone the molar amount of the o-dihydroxybenzene in said mixture being larger than the molar amount of the 3-pyrazolidinone, and the p-dihydroxybenzene if any being present in a molar ratio of at most 5 % with respect to the o-dihydroxybenzene can be used. Other type of developing agents suitable for use in accordance with the present invention are reductones e.g. ascorbic acid derivatives.

The developing agent or a mixture of developing agents can be present in an alkaline processing solution, in the photographic material or the image receiving material. In case the developing agent or a mixture of developing agents is contained in the photographic material and/or image receiving material, the processing solution can be merely an aqueous alkaline solution that initiates and activates the development.

In the DTR process the photographic element is developed in the

WO 95/16224

presence of a silver halide solvent. Preferably used silver halide solvents are water soluble thiosulphate compounds such as ammonium and sodium thiosulphate, or ammonium and alkali metal thiocyanates. Other useful silver halide solvents (or "complexing agents") are described in the book "The Theory of the Photographic Process" edited by T.H. James, 4th edition, p. 474-475 (1977), in particular sulphites and uracil. Further interesting silver halide complexing agents are cyclic imides, preferably combined with alkanolamines, as described in US-P 4,297,430 and US-P 4,355,090. 2-mercaptobenzoic acid derivatives are described as silver halide solvents in US-P 4,297,429, preferably combined with alkanolamines or with cyclic imides and alkanolamines. Dialkylmethylenedisulfones can also be used as silver halide solvent.

18

PCT/EP94/04095

The silver halide solvent is preferably present in the processing solution but may also be present in one or more layers comprised on the support of the imaging element and/or receiving material. When the silver halide solvent is incorporated in the photographic material it may be incorporated as a silver halide solvent precursor as disclosed in e.g. Japanese published unexamined patent applications no. 15247/59 and 271345/63, US-P 4,693,955 and US-P 3,685,991.

The processing solution for use in the production of black-and-white photographs in opaque security documents according to the present invention may comprise other additives such as e.g. thickeners, preservatives, detergents e.g. acetylenic detergents such as SURFYNOL 104, SURFYNOL 465, SURFYNOL 440 etc. all available from Air Reduction Chemical Company, New York.

The DTR-process is normally carried out at a temperature in the range of  $10^{\circ}\text{C}$  to  $35^{\circ}\text{C}$ .

Further details about the black-and-white DTR process and also about the dye diffusion transfer process and image receiving materials used therein are described in Research Disclosure November 1976, item 15162.

The present invention will now be illustrated by the following examples without however limiting it thereto. All ratios, percentages and parts are by weight unless otherwise specified.

19

#### EXAMPLE 1

- Preparation of photographic element for use in the DTR process

A gelatino silver halide emulsion was prepared by slowly running with stirring an aqueous solution of 1 mole of silver nitrate per liter into a gelatine solution containing per mole of silver nitrate 41 g of gelatin, 1.2 mole of sodium chloride, 0.08 mole of potassium bromide and 0.01 mole of potassium iodide.

The temperature during precipitation and the subsequent ripening process lasting three hours was kept at 40°C.

Before cooling, shredding and washing 214 g of gelatin were added per mole of silver halide. The washed noodles were molten and another 476 g of gelatin were added per mole of silver halide during the chemical ripening. After ripening 285 g of gelatin in the form of a 20 % aqueous solution were added to the emulsion per mole of silver halide as well as hydroquinone in an amount such that after coating 0.9 g of hydroquinone were present per m<sup>2</sup> and 1-phenyl-4,4dimethyl-3-pyrazolidinone in an amount such that 0.21 g thereof were present per m<sup>2</sup>. The emulsion was coated at one side of a subbed water-resistant paper support consisting of a paper having a weight of 110  $g/m^2$  coated at both sides with a polyethylene stratum at a ratio of 20  $g/m^2$  per side.

The emulsion was coated in such a way that an amount of silver equivalent to 1.5 g of silver nitrate was applied per m<sup>2</sup>. The amount of gelatin corresponding therewith is  $8.93~\mathrm{g/m^2}$  since the gelatin to silver nitrate weight ratio was 5.97.

- Preparation of image receiving material for use in the DTR process

One side of a paper support having a weight of  $100 \text{ g/m}^2$  being coated at both sides with a polyethylene layer of 20  $\mathrm{g/m^2}$  was coated after corona treatment at a dry coverage of 2.5  $g/m^2$  of gelatin and 1.3  $g/m^2$  of interference pigment from the following coating composition :

carboxymethyl cellulose	12	g
gelatin	38.5	g
3 % aqueous dispersion of colloidal Ag2S.NiS nuclei	14	ml

20

4 % aqueous solution of formaldehyde 12 ml aqueous dispersion of blue PALIOSECURE type EC 1408 (tradename) containing 30 % of said blue pigment and 8 % of gelatin 80 g 12.5 % solution of saponine in ethanol/water (20/80) 20 ml

- Printing of the image receiving material with pattern of interference pigment

In the area lateron to be DTR-processed to form therein a portrait of a person being owner of a passport, a guilloche pattern was gravure-printed using an ink having the following composition:

15 % dispersion of green PALIOSECURE type EC 1407 (tradename) in a cellulose nitrate solution in liquid polyethylene glycol ether.

The printed ink pattern was dried at the air.

- Printing of the image receiving material with pattern of graphical and numerical information using a blue non-iridiscent ink

The printing of said information was carried out in the background area having a blue color due to the presence of said interference pigment PALIOSECURE type EC 1408 (tradename). The printing of the blue non-iridescent ink proceeded at such density that there was no visible contrast at 90° inspection angle (being also the usual copying angle) between the background and the thus printed area.

DTR-image formation

The above defined photographic element was image-wise exposed in a reflex camera to obtain therein in the guilloche covered area a photograph (portrait) of the passport owner.

The photo-exposed element was pre-moistened with a processing liquid as defined hereinafter.

The contact time of the photo-exposed element with said liquid was 6 seconds before being pressed together with the image-receiving material as defined above. The transfer processor employed was a COPYPROOF (registered trade name of AGFA-GEVAERT N.V.) type CP 380. The transfer contact time was 30 seconds. In the image-receiving layer a positive black-and-white (silver image) portrait of the photographed person was obtained.

21

WO 95/16224 PCT/EP94/04095

## - Composition of the processing liquid:

hydroxyethyl cellulose	1.0 g
Ethylenediaminetetraacetic acid tetrasodium salt	2.0 g
$Na_2SO_3$	45.0 g
$Na_2S_2O_3$	14.0 g
KBr	` 0.5 g
1-Pheny1-5-mercapto-tetrazole	0.1 g
1-(3,4-Dichlorophenyl)-1H-tetrazole-5-thiol	0.02 g
N-methyl-ethanolamine	45.0 ml
N-methyl-diethanolamine	30.0 ml
Water up to	1 1

When viewed under an angle of 90° in daylight the color of the non-printed area around the portrait was blue due the presence of uniformly distributed therein blue PALIOSECURE type EC 1408 (tradename). In the portrait area containing green PALIOSECURE type EC 1407 (tradename) black silver was missing and the guilloche pattern had a bluish-green color.

When viewing the document under an angle less than 45° the background area containing blue PALIOSECURE type EC 1408 (tradename) changed to become yellowish. Under that angle the graphical information and numerical information printed with said blue non-iridescent ink became legible.

On copying the obtained document with a color copier (CANON CLC 500) the above defined color changes were not reproduced in the obtained print. The guilloche pattern covering the portrait area can hardly be seen in the obtained print, and the blue printed graphical information is not copied.

#### EXAMPLE 2

- Preparation of image-receiving element for use in dye diffusion transfer process

An opaque polyvinyl chloride sheet having a thickness of 0.200 mm was after corona treatment coated at one side with the following compositions for forming a subbing layer and mordanting layer respectively:

## 1. Subbing layer coating composition

gelatin	4 g	j >
aqueous dispersion of blue PALIOSECURE type EC 1408 (trade:	name)	•
containing 30 % of said blue pigment and 8 % of gelatin	200	g
ingredient A 40 % solution dispersed in aqueous medium	250	m1
5 % solution of siloxane compound in ethanol	125	m1
12.5 % solution of saponine in ethanol/water 20/80	20	m1

Ingredient A is a polyester-polyurethane having the same chemical composition as described in US-P 4,902,593, column 2, lines 64-68 and column 3, lines 1-8.

The coating composition was applied coated at a dry coverage of  $0.4~\rm g/m^2$  of gelatin and  $1.2~\rm g.m^2$  of interference pigment.

#### 2. Coating composition of the mordanting layer

gelatin	20	g
mordant M (20 % solution)	250	m1
saponine (12 %) and wetting agent W (5 %) in water	32	m1
aqueous 4 % solution of formaldehyde	10	m1

Mordant M on the basis of an epoxidized cationic polymer has the same composition as described in US-P 4,902,593, column 7, lines 14-42.

The coating composition was applied at a dry coverage of 0.9  $\rm g/m^2$  of gelatin.

The above defined image-receiving material was processed in combination with a photographic dye diffusion transfer material as described in the Example of U.S. Pat. No. 4,496,645, which material was exposed to reproduce thereon a portrait. The exposed material was kept for 1 minute in contact with the above defined image-receiving material after being led through a diffusion transfer apparatus COPYPROOF CP 38 (tradename of Agfa-Gevaert N.V. Belgium) having in its tray the following basic processing liquid:

sodium hydroxide	25 g
sodium orthophosphate	25 g
cyclohexane dimethanol	25 g
2.2' methylpropylpropane diol	25 g
N-ethylbenzene-pyridinium chloride	0.5 g
distilled water up to	1000 ml

After leaving the processing tray the image-receiving sheet was led through a second tray containing an aqueous 5 % solution of the already mentioned wetting agent W corresponding with the following formula: iso-nonyl-phenoxy- $(CH_2-CH_2-O)_9$ -H and potassium iodide (ref. EP 0250657).

After drying the processed sheet material it was laminated as described in US-P 4,902,593 to obtain a sealed I.D. card.

#### EXAMPLE 3

According to Example 3 the interference pigments were applied uniformly in a mordanting layer of an image-receiving material suited for use in a dye diffusion transfer process.

#### - Preparation of the image-receiving element

An opaque polyvinyl chloride sheet having a thickness of 0.200 mm was after corona treatment coated at one side with the following compositions for forming a subbing layer and mordanting layer respectively:

## 1. Coating composition of the subbing layer

gelatin	20 g
ingredient A 40 % solution dispersed in aqueous medium	250 ml
5 % solution of siloxane compound in ethanol	125 ml
12.5 % solution of saponine in ethanol/water 20/80	20 ml

The coating composition was applied coated at a dry coverage of 0.4  ${\rm g/m^2}$  of gelatin.

# 2. Coating composition of the mordanting layer

gelatin	12 g
aqueous dispersion of blue PALIOSECURE type EC 1408 (trade	ename)
containing 30 % of said blue pigment and 8 % of gelatin	100 g
mordant M (20 % solution)	250 ml
saponine (12 %) and wetting agent W (5 %) in water	32 ml
aqueous 4 % solution of formaldehyde	10 ml

The coating composition was applied at a dry coverage of 0.9  $\rm g/m^2$  of gelatin, and 1.3  $\rm g/m^2$  of interference pigment.

#### EXAMPLE 4

Example 3 was repeated with the difference that the interference pigments were applied uniformly in a gelatin top coat covering the mordanting layer. The dried top coat contained 0.5 g/m $^2$  of gelatin and 1.3 g/m $^2$  of interference pigment.

25

#### CLAIMS

- 1. A document containing a photograph obtained by the silver complex or dye diffusion transfer process in an image receiving layer containing light interference pigments uniformly distributed in said image receiving layer, and/or in a layer overlaying or underlying said image receiving layer, said layer(s) being present on an opaque support, characterised in that said interference pigments are composed of silicate platelets coated with either:
  - A) a first layer of highly refractive metal oxide, and
  - B) a second black layer essentially consisting of metallic iron, molybdenum and/or wolfram, or coated with:
  - A) a first layer of highly refractive metal oxide, and
  - B) a second black layer essentially consisting of carbon or metal, and
  - C) a third layer of highly refractive metal oxide.
- 2. Document according to claim 1, wherein in said document said uniformly distributed interference pigments of a particular color are present in combination with a printed pattern containing interference pigments of a color different from the color of the uniformly distributed interference pigments when viewed under the angle of reflection under which photocopying could take place.
- 3. Document according to claim 1 or 2, wherein in said document said uniformly distributed interference pigments are present in conjunction with different and pattern-wise printed interference pigments on a background having a color substantially the same as the color reflected by said pattern-wise printed interference pigments under the angle of reflection under which photocopying could take place.
- 4. Document according to any of claims 1 to 3, wherein in said document said uniformly distributed interference pigments are present in conjunction with uniformly or pattern-wise printed colorants (colored pigments or dyes), and/or white light-diffusing pigments.
- 5. Document according to any of claims 1 to 4. wherein said photograph is formed by the silver salt diffusion transfer process

26

in an image-receiving layer comprising physical development nuclei.

- 6. Document according to any of claims 1 to 4, wherein said photograph is formed by a dye diffusion transfer process in an image-receiving layer containing a mordant for a dye transferred from an image-wise exposed and developed silver halide emulsion material.
- 7. Document according to any of claims 1 to 6, wherein said pigments in the form of platelets have a largest surface diameter between 5 and 200  $\mu m\,.$
- 8. Document according to any of claims 1 to 7, wherein the thickness of said interference pigments is between 0.1  $\mu m$  and 0.6  $\mu m$  .
- 9. Document according to any of the preceding claims, wherein said uniformly distributed interference pigments are present in said document in a coverage of 0.3  $\rm g/m^2$  to 10  $\rm g/m^2$ .

## INTERNATIONAL SEARCH REPORT

International Application No PCT/L, 94/04095

A. CLASSIFICATION OF SUBJECT MATTER IPC 6 G03C5/08 B41M3/14 G03C8/52 G03C8/28 B42D15/10 G03C5/08 IPC 6 According to International Patent Classification (IPC) or to both national classification and IPC **B. FIELDS SEARCHED** Minimum documentation searched (classification system followed by classification symbols) G03C B41M **B42D** IPC 6 Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Electronic data base consulted during the international search (name of data base and, where practical, search terms used) C. DOCUMENTS CONSIDERED TO BE RELEVANT Relevant to claim No. Citation of document, with indication, where appropriate, of the relevant passages Category 1,5,6 EP,A,0 447 692 (AGFA-GEVAERT NV) 25 Y September 1991 see page 9, line 45 - line 52 see page 5, line 5 - line 9 1,5 EP,A,O 490 416 (AGFA-GEVAERT NV) 17 June Y 1992 see page 11, line 18 - line 19 1,5,6 CA,A,2 123 783 (BASF AG) 24 June 1993 Y cited in the application see page 11, line 30 - line 32 see page 1, line 28 - page 2, line 5 US,A,4 151 666 (THOMAS RAPHAEL ET AL) 1 May 1979 cited in the application see column 3, line 6 - line 17 -/--Patent family members are listed in annex. Further documents are listed in the continuation of box C. "T" later document published after the international filing date Special categories of cited documents: or priority date and not in conflict with the application but cited to understand the principle or theory underlying the "A" document defining the general state of the art which is not considered to be of particular relevance invention "X" document of particular relevance; the claimed invention "E" earlier document but published on or after the international cannot be considered novel or cannot be considered to filing date involve an inventive step when the document is taken alone "L" document which may throw doubts on priority claim(s) or "Y" document of particular relevance; the claimed invention which is cited to establish the publication date of another cannot be considered to involve an inventive step when the citation or other special reason (as specified) document is combined with one or more other such docu-"O" document referring to an oral disclosure, use, exhibition or ments, such combination being obvious to a person skilled other means in the art. "P" document published prior to the international filing date but "&" document member of the same patent family later than the priority date claimed Date of mailing of the international search report Date of the actual completion of the international search 2 1. 04. 95 30 March 1995 Authorized officer Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Bolger, W Fax: (+31-70) 340-3016

. 1

# INTERNATIONAL SEARCH REPORT

Internation 1 Application No
PCT/Er 94/04095

		PCT/Er 94/04095	
(Continua	tion) DOCUMENTS CONSIDERED TO BE RELEVANT	Relevant to	claim No.
ategory *	Citation of document, with indication, where appropriate, of the relevant passages		
	US, A, 4 428 997 (JOSEPH SHULMAN) 31 January 1984	3,	4
	see column 3, line 35 - column 5, line 55  NL,A,8 901 256 (JOH.ENSCHEDE EN ZONEN  1990	3,	,4
•	GRAFISCHE INRICHTING BV) 17 December 1990 see page 4, line 2 - line 4; claim 1		
	FR,A,2 429 292 (ARJOMARI-PRIOUX) 18 January 1980 see claim 1	1	
A	EP,A,O 317 514 (LIPATEC ETABLISSEMENT) 24 May 1989	1	
	cited in the application see claim 1		
Ρ,Α	EP,A,0 588 407 (AGFA-GEVAERT NV) 23 March 1994	1	
	see page 3, line 23 - line 37		
1			

# INTERNATIONAL SEARCH REPORT

In...mation on patent family members

Internatic 'Application No
PCT/Er 94/04095

Patent document sited in search report	Publication date	Patent family member(s)		Publication date
EP-A-0447692	25-09-91	JP-A- US-A-	4261843 5213648	17-09-92 25-05-93
EP-A-0490416	17-06-92	JP-A- US-A-	4333048 5194347	20-11-92 16-03-93
CA-A-2123783	24-06-93	DE-A- WO-A- EP-A-	4141069 9312182 0616626	17-06-93 24-06-93 28-09-94
US-A-4151666	01-05-79	NONE		
US-A-4428997	31-01-84	NONE		
NL-A-8901256	17-12-90	NONE		
FR-A-2429292	18-01-80	NONE		
EP-A-0317514	24-05-89	CH-A- AU-A- DE-A- JP-A- US-A-	672687 2569788 3873034 1155381 5087507	15-12-89 25-05-89 27-08-92 19-06-89 11-02-92
EP-A-0588407	23-03-94	JP-A- US-A-	6194804 5340692	15-07-94 23-08-94